

**K092073 MONOPOLY PEDICLE SCREW SYSTEM**Aug 7, 2009  
30 days to decisionK092073 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k092073/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jul 8, 2009
Decision date	Aug 7, 2009
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Signus Medical</b>
Location	Washington, DC, US
Contact	THOMAS HOGHAUG
510(k) history	2 submissions · 2 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092073/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 19, 2026